AMENDMENT TO THE CLAIMS

- 1. (Currently Amended) A method of determining whether a <u>human</u> test subject has, is at risk of developing, or <u>has an increased likelihood</u> may have or be at risk of developing a titin-related disease or condition of the heart, said method comprising obtaining a sample from said test subject and analyzing a nucleic acid molecule of said sample to determine whether the test subject has a mutation in a naturally occurring <u>human</u> *titin* gene, wherein the presence of said mutation is an indication that said test subject has, is at risk of developing, or <u>has an increased</u> <u>likelihood</u> may have or be at risk of developing a titin-related disease or condition of the heart.
- 2. (Previously Presented) The method of claim 1, wherein said analyzing of said nucleic acid molecule comprises using nucleic acid molecule primers specific for the *titin* gene for nucleic acid molecule amplification of the *titin* gene by the polymerase chain reaction.
- 3. (Previously Presented) The method of claim 1, wherein said analyzing of said nucleic acid molecule comprises sequencing *titin* nucleic acid molecules from said test subject.
 - 4-5. (Canceled).
 - 6. (Original) The method of claim 1, wherein said disease or condition is heart failure.
 - 7. (Canceled).

- 8. (Withdrawn) A method for identifying a compound that can be used to treat or to prevent heart failure, said method comprising contacting an organism comprising a *titin* mutation and having a phenotype characteristic of heart failure with said compound, and determining the effect of said compound on said phenotype, wherein detection of an improvement in said phenotype indicates the identification of a compound that can be used to treat or to prevent heart failure.
 - 9. (Withdrawn) The method of claim 8, wherein said organism is a zebrafish.
- 10. (Withdrawn) The method of claim 8, wherein said *titin* mutation is the *pickwick* mutation.
- 11. (Withdrawn) A method of treating or preventing heart failure in a patient, said method comprising administering to said patient a compound identified using the method of claim 8.
- 12. (Withdrawn) The method of claim 11, wherein said patient has a mutation in the *titin* gene.
- 13. (Withdrawn) The method of claim 12, wherein said mutation is the *pickwick* mutation.

- 14. (Withdrawn) A non-human animal comprising a mutation in a titin gene.
- 15. (Withdrawn) The non-human animal of claim 14, wherein the non-human animal is a zebrafish.
- 16. (Withdrawn) The non-human animal of claim 14, wherein the mutation is in a cardiac-specific exon of said *titin* gene.
- 17. (Withdrawn) The non-human animal of claim 16, wherein the mutation is in the N2B exon of said *titin* gene.
- 18. (Withdrawn) The non-human animal of claim 14, wherein the mutation results in the presence of a stop codon in said *titin* gene.
- 19. (Withdrawn) The non-human animal of claim 14, wherein the mutation is the *pickwick* mutation.
- 20. (Currently Amended) The method of claim 1, wherein the test subject <u>has an</u> increased likelihood may have or be at risk of developing a titin-related disease or condition.
- 21. (Previously Presented) The method of claim 1, wherein the mutation is in a cardiacspecific exon of said *titin* gene.

- 22. (Previously Presented) The method of claim 1, wherein the mutation is in the N2B exon of said *titin* gene.
- 23. (Previously Presented) A method of determining whether a test subject has an increased likelihood of a titin-related disease or condition of the heart, or facilitating determination of the etiology of an existing heart condition, the method comprising obtaining a sample from the test subject and analyzing a nucleic acid molecule of the sample to determine whether the test subject has a mutation in a *titin* gene, wherein the presence of a mutation in a *titin* gene is an indication that the test subject has an increased likelihood of a titin-related disease or condition of the heart, or provides information as to the etiology of an existing heart disease or condition.
- 24. (Previously Presented) The method of claim 23, wherein the method is carried out to determine whether the test subject has an increased likelihood of a titin-related disease or condition of the heart.
- 25. (Previously Presented) The method of claim 23, wherein the method is carried out to facilitate determination of the etiology of an existing heart disease or condition in the test subject.
- 26. (Previously Presented) The method of claim 23, wherein said analyzing of said nucleic acid molecule comprises using nucleic acid molecule primers specific for the *titin* gene

for nucleic acid molecule amplification of the titin gene by the polymerase chain reaction.

- 27. (Previously Presented) The method of claim 23, wherein said analyzing of said nucleic acid molecule comprises sequencing *titin* nucleic acid molecules from said test subject.
- 28. (Previously Presented) The method of claim 23, wherein said test subject is a mammal.
 - 29. (Previously Presented) The method of claim 23, wherein said test subject is human.
- 30. (Previously Presented) The method of claim 23, wherein said disease or condition is heart failure.
- 31. (Previously Presented) The method of claim 23, wherein the mutation is in a cardiacspecific exon of said *titin* gene.
- 32. (Previously Presented) The method of claim 23, wherein the mutation is in the N2B exon of said *titin* gene.